



*Institute of Paper Science and Technology
Atlanta, Georgia*

IPST Technical Paper Series Number 821

The Practical Influence of Heterogeneity on Accelerated Creep in Paper

D. Coffin and C. Habeger

November 1999

Submitted to
Tappi Journal

Copyright® 1999 by the Institute of Paper Science and Technology

For Members Only

INSTITUTE OF PAPER SCIENCE AND TECHNOLOGY PURPOSE AND MISSIONS

The Institute of Paper Science and Technology is an independent graduate school, research organization, and information center for science and technology mainly concerned with manufacture and uses of pulp, paper, paperboard, and other forest products and byproducts. Established in 1929, the Institute provides research and information services to the wood, fiber, and allied industries in a unique partnership between education and business. The Institute is supported by 52 North American companies. The purpose of the Institute is fulfilled through four missions, which are:

- to provide a multidisciplinary education to students who advance the science and technology of the industry and who rise into leadership positions within the industry;
- to conduct and foster research that creates knowledge to satisfy the technological needs of the industry;
- to serve as a key global resource for the acquisition, assessment, and dissemination of industry information, providing critically important information to decision-makers at all levels of the industry; and
- to aggressively seek out technological opportunities and facilitate the transfer and implementation of those technologies in collaboration with industry partners.

ACCREDITATION

The Institute of Paper Science and Technology is accredited by the Commission on Colleges of the Southern Association of Colleges and Schools to award the Master of Science and Doctor of Philosophy degrees.

NOTICE AND DISCLAIMER

The Institute of Paper Science and Technology (IPST) has provided a high standard of professional service and has put forth its best efforts within the time and funds available for this project. The information and conclusions are advisory and are intended only for internal use by any company who may receive this report. Each company must decide for itself the best approach to solving any problems it may have and how, or whether, this reported information should be considered in its approach.

IPST does not recommend particular products, procedures, materials, or service. These are included only in the interest of completeness within a laboratory context and budgetary constraint. Actual products, procedures, materials, and services used may differ and are peculiar to the operations of each company.

In no event shall IPST or its employees and agents have any obligation or liability for damages including, but not limited to, consequential damages arising out of or in connection with any company's use of or inability to use the reported information. IPST provides no warranty or guaranty of results.

The Institute of Paper Science and Technology assures equal opportunity to all qualified persons without regard to race, color, religion, sex, national origin, age, disability, marital status, or Vietnam era veterans status in the admission to, participation in, treatment of, or employment in the programs and activities which the Institute operates.

The Practical Influence of Heterogeneity on Accelerated Creep in Paper

Douglas Coffin and Chuck Habeger

ABSTRACT

According to theoretical work done by the authors (5,6), sheet heterogeneity can affect accelerated creep in paper. Experiments were conducted to see whether these influences have any impact on everyday papermaking decisions. We report that papers made from furnishes of mixed fiber type do not experience additional accelerated creep, whereas sheets formed from plies of different hygroexpansivity exhibit extra accelerated creep. An attempt to create fiber microcompressions by Hobart mixer treatment did not change the accelerated creep performance.

Application:

Papermakers should be aware that multi-ply paperboard products could fail sooner than expected under load in cyclic humidity environments. A similar concern is not raised for paper made from a blend of different furnishes.

Accelerated creep is an established phenomenon in wood (1), paper (2), and other materials (3,4). Hydrophilic materials generally creep faster at a high relative humidity (RH) than at a low RH. However, under the proper circumstances, creep can be greater in an environment that is continually cycled between low and high RH levels than when it is held at the high humidity extreme. When this is observed, the material is said to experience accelerated creep. Figure 1 presents a typical experiment exhibiting accelerated creep in paper. There, we have plotted creep strain versus log time. The sample was maintained under tensile load for three hours at 80% RH. Then, keeping the load constant, the humidity was cycled at one-hour intervals between 80 and 30% RH. Notice that the creep under cyclic conditions is far above the time-projected wet creep curve.

The driving mechanism for accelerated creep is a contentious subject (5). We advocate (5,6) a cyclic moisture-induced cyclic stress explanation (3,5,6,7). We demonstrate that those materials, which experience accelerated creep, show more creep in cyclic load testing than when they are held at a constant, average load. We argue that accelerated creep is the result of this nonlinear creep constitutive behavior and the localized load cycling produced in cyclic humidity environments. Sorption-induced moisture gradients are a likely source of load cycling that is caused by moisture cycling. In our view, sheet heterogeneity in response to moisture is another possibility. We undertook the experimental work reported below in order to see if "heterogeneity-driven accelerated creep" is of practical interest to papermakers.

To envision the heterogeneity type of accelerated creep, imagine adjacent structural elements in the paper that are parallel to the direction of the applied load. If these elements have different moisture sensitivities, their stresses will be redistributed during sorption, and moisture cycling will lead to out-of-phase load cycling of each

element and in turn to extra creep. If this extra creep overcompensates for the low creep rates at low moisture content, a sample in a cycling moisture environment can creep more than another sample held at the highest moisture content (5,6). As an example, consider a sheet made of two separate plies under a constant tensile load. Let the hygroexpansion of one of the plies be greater than the other ply. As the sheet absorbs moisture, one side attempts to expand more than the other. Part of the tensile load is shifted to the less hygroexpansive ply. Upon desorption, the reverse occurs: the more moisture sensitive ply carries most of the tensile load. During moisture cycling, the loads on each of the two plies cycle out-of-phase. This same kind of action might also take place, but on a fiber level, if fibers of differing hygroexpansivity are bonded together. Wood fibers expand at least an order of magnitude more laterally than axially with the uptake of water (8); thus identical fibers aligned crosswise will also cycle their load sharing in a cyclic humidity chamber. In a like manner, heterogeneity in the moisture dependence of stiffness and creep compliance can also generate load cycling and accelerated creep.

We just identified three types of heterogeneity (through-sheet heterogeneity, fiber-to-fiber heterogeneity, and fiber-level heterogeneity due to fiber anisotropy) as candidates for initiating load cycling and accelerated creep. We decided to conduct experiments that would shed light on the importance of these mechanisms and on the degree that papermakers can influence accelerated creep by altering heterogeneity parameters. To do this, we procured furnishes that produced papers which responded differently to changing relative humidity. We formed blended sheets and multi-ply sheets of furnishes composed of fibers of different hygroexpansivity and compared their accelerated creep performance to that of sheets made of a single fiber type.

We wanted to work with common fibers having significant differences in hygroexpansion. Although drying restraint has a major influence on hygroexpansion (9), we cannot expect different sections of our samples to have different drying histories; therefore, we could not reasonably select drying restraint as a variable for our accelerated creep experiments. Hygroexpansion, defined as change in percent length divided by change in percent moisture (10), depends on furnish (11). Mechanical pulps shrink less than low yield chemical pulps during drying, and the dimensional stability of mechanical pulps does not benefit as much from restrained drying (9). Fully restrain-dried TMP sheets are about 10% more hygroexpansive than fully restrained bleach kraft sheets (9). In our accelerated creep testing, we are changing relative humidity; therefore, we are interested in the change in length induced by changes in relative humidity. Sheets made from mechanical pulps have about a 10% greater rate of change in moisture with RH than sheets made of chemical pulps. Therefore, we concluded that fully restrained sheets formed from mechanical and chemical pulps would be different in their RH-based hygroexpansions, and that these differences would be representative of those encountered in papermaking. We investigated the influence of through-sheet heterogeneity by testing samples made by wet pressing plies of different furnish combinations and then drying them under restraint. Fiber-to-fiber heterogeneity was assessed by comparing the performances of blended furnishes with those of pure furnishes.

We have argued (5,6) that the anisotropy in a wood fiber's response to moisture leads to a fiber-level heterogeneity in the sheet that contributes to paper accelerated creep. We experimentally address the influence of fiber anisotropy by looking at the accelerated creep of sheets made of fibers treated to be less anisotropic. Wood fibers are filament-wound structures of crystalline cellulose microfibrils held together by a lignin-hemicellulose matrix. They are stiff and moisture insensitive in the axial direction and compliant and hygroexpansive radially. High-consistency mixing curls and microcompresses wood fibers (12). The curl, but not the microcompressions, can be removed by subsequent low consistency treatment. The microfibrillar structure is buckled and folded at the microcompressions. This preferentially reduces axial stiffness and increases axial moisture sensitivity. As suggested by Derek Page, we used a Hobart kitchen mixer (13) as a means to induce microcompressions and reduce fiber anisotropy.

SAMPLES

We chose a newsprint TMP and a bleached softwood kraft pulp as the test furnishes for our blend-ply experiments. These furnishes were received as never-dried pulps, and we did no refining. After removing the fines, the freenesses were measured as 721 CSF for the BKS and as 767 CSF for the TMP. Sheets of 60 g/m² were formed on a Noble and Wood 8-inch square handsheet mold. The sheets were wet pressed (singly and in stacks of four) at a common pressure of 50 psi for 5 minutes. All sheets were restrained-dried on a static drum drier. Single-ply handsheets were made from the TMP furnish (sheets A), the BKS furnish (sheets B), and a 50-50 retained-on-the-wire blend of the TMP and BKS fibers (sheets AB). The D sheets were formed from the BKS pulp after it was processed in the Hobart mixer at 20% consistency for two hours. The pulp was then processed in a British Disintegrator at 1.2% consistency to remove curl. Four-ply handsheets were formed with TMP plies (sheets AAAA), BKS plies (sheets BBBB), two TMP plies in the middle and a BKS ply on each side (sheets BAAB), and two BKS plies in the middle and a TMP ply on each side (sheets ABBA).

We used our variable-humidity creep tester (5,6) to measure hygroexpansion of the samples between 30 and 80% RH. We applied a tensile load of 100 grams (400 grams for four-ply samples). This was large enough to straighten the samples, but it was insufficient to cause appreciable elastic or creep strain. The 100-gram load supplies only about 5% of the ultimate tensile load of the weakest sample. After conditioning with several humidity cycles, we recorded the strain difference between 80 and 30% RH as the hygroexpansion. With this technique, the standard deviations in the hygroexpansions were about .01%. Pertinent average physical properties are provided in Table I.

Table I
Physical Properties of Samples

Sheet Type	Density	Basis Weight	Tensile Strength	Stretch	Long. Sonic Modulus	Hygroexpansion 30-80% RH
	g/cm ²	g/m ²	kN/m	%	km ² /s ²	%
A	.314	61.9	1.84	1.6	4.98	0.39
B	.317	61.9	0.68	1.1	3.95	0.32
AB	.305	59.6	1.41	2.0	4.81	0.36
D	.491	60.3	1.69	2.4	6.79	0.35
AAAA	.332	254	8.41	2.7	5.23	0.34
BBBB	.356	250	3.36	1.9	4.84	0.30
ABBA	.333	256	5.58	2.2	4.94	0.32
BAAB	.347	258	5.81	2.5	4.73	0.33

ACCELERATED CREEP TESTING

We did the accelerated creep testing in our controlled-environment, tensile creep tester as before (5,6). Samples were mounted in the chamber and equilibrated to 80% RH. The tensile load was applied and held for three hours, at which time the RH was cycled between 30 and 80% RH at one-hour intervals. The sample lengths were monitored, and creep strain was determined as a function of time. We made from two to four replications (generally three) of each test and averaged the results.

Notice that there is a considerable difference in tensile strength between samples. In previous testing, we observed that accelerated creep depends on tensile load. Accelerated creep measurements could be compared at the same nominal load or at the same portion of tensile strength. Each approach has its advantages, so we decided, in all cases, to do it both ways.

The accelerated creep numbers for A, B, and AB at 25% of the sample's 50% RH tensile strength and at 25% of the 50% RH strength of sample B are presented in Table II. The standard deviations are listed in parentheses. We define accelerated creep as the ratio of the creep versus log time slope taken after and before humidity cycling begins (see Figure 1).

Table II
Blend Accelerated Creep Results

Sheet Type	Accelerated Creep at 25% of Sample's Tensile Strength	Accelerated Creep at 25% of Sample B's Tensile Strength
A	5.16 (.3)	8.30 (.4)
B	4.48 (.1)	4.48 (.1)
AB	4.61 (.1)	6.00 (.3)

As in the blend testing, we made multi-ply accelerated creep comparisons at the same load and at the same portion of tensile strength. This time, we did equal-load testing at two levels: 25% and 35% of the tensile strength of BBBB at 50% RH. The results are listed in Table III.

Table III
Multi-ply Accelerated Creep Results

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of sample BBBB's tensile strength	Accelerated Creep at 35% of sample BBBB's tensile strength
AAAA	3.79 (.1)	4.21 (.1)	5.21 (.1)
BBBB	3.89 (.1)	3.89 (.1)	3.26
ABBA	4.51 (.2)	5.24 (.1)	4.12
BAAB	4.17 (.2)	4.62 (.1)	4.30

The comparisons between accelerated creep with and without treatment in the Hobart mixer follow as Table IV.

Table IV
Treated Accelerated Creep Results

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of B's tensile strength
B	4.48 (.1)	4.48 (.1)
D	3.71 (.1)	4.51 (.2)

We adopted the ratio of the log time creep slopes as our parameter to quantify accelerated creep (4,5,6) before we seriously began comparing the values of different samples. In truth, this decision is arbitrary and warrants reconsideration. First let us look more closely at the present method. Figure 1 is a representative accelerated creep plot. It is our custom to quantify accelerated creep as the ratio of the slope of the line tangent to the local maximums of the cyclic creep curve to the slope of the constant humidity creep curve in the early part of the experiment. Notice that the construction in the cyclic humidity portion of the experiment forms an almost straight line, whereas the slope of the creep in the initial high-humidity regime is not constant. That is, at these levels of loads and humidity cycling, creep is log linear under cyclic humidity, but it is not in the initial wet state. Therefore, the process of slope determination is not obvious for the wet state. We have arbitrarily taken it as the tangent to the creep line just prior to the beginning of humidity cycling. Other materials, such as Kevlar fibers (4,5,6), show log time linear creep in both regimes, and this determination of accelerated creep is more consistent.

At relatively small loads and short times, creep in paper follows the power law; it only becomes log linear at long times or high loads (14) or, as we have demonstrated,

under cyclic humidity. Power law creep forms a straight line when log creep is plotted versus log time. The same data as in Figure 1 is plotted in Figure 2; however, the logarithm of creep strain is now on the y-axis. This time, things are reversed: the wet-state creep is linear, whereas the cyclic humidity portion is not linear. We could have just as logically used the log-log plot to define accelerated creep. For paper, we see no overwhelming reason to prefer one method above the other.

Notice, from Figure 1 or Figure 2, that a significant portion of the extra elongation induced by moisture cycling comes during the first rewetting cycle. Both of these accelerated creep calculations ignore this part of the creep. They take the cyclic humidity slope between peaks, after the first rewetting. This is arguably inappropriate. For paper, creep rates fall as creep increases. A sample that has a large response upon first rewetting will have a smaller subsequent creep rate. A calculation that doesn't take this into account could misrepresent relative accelerated creep rates. A third alternative is to use the slope of the line between the beginning and the end of the cyclic humidity strain of the log-log plot as the numerator in the accelerated creep calculation (see Figure 2). This slope clearly depends on the length of the test. However, it does account for all the first rewetting creep, and we used a common humidity cycling history for all tests.

The alternative accelerated creep calculations follow in the tables below. Tables with "a" designation result from the slopes on the log-log plots. Those with "b" designations use the full creep slope in the numerator.

Table II-a
Blend Accelerated Creep Results
log-log plot

Sheet Type	Accelerated Creep at 25% of Sample's Tensile Strength	Accelerated Creep at 25% of Sample B's Tensile Strength
A	1.50	1.55
B	1.45	1.45
AB	1.55	1.51

Table II-b
Blend Accelerated Creep Results
full log-log plot

Sheet Type	Accelerated Creep at 25% of Sample's Tensile Strength	Accelerated Creep at 25% of Sample B's Tensile Strength
A	2.37	2.86
B	2.26	2.26
AB	2.34	2.52

Table III-a
Multi-ply Accelerated Creep Results
log-log plot

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of sample BBBB's tensile strength	Accelerated Creep at 35% of sample BBBB's tensile strength
AAAA	1.56	1.18	1.24
BBBB	1.40	1.40	1.51
ABBA	1.40	1.44	1.50
BAAB	1.65	2.32	1.46

Table III-b
Multi-ply Accelerated Creep Results
full log-log plot

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of sample BBBB's tensile strength	Accelerated Creep at 35% of sample BBBB's tensile strength
AAAA	1.85	1.96	2.03
BBBB	2.13	2.13	2.08
ABBA	2.07	2.34	2.27
BAAB	2.38	3.44	2.38

Table IV-a
Treated Accelerated Creep Results
log-log plot

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of B's tensile strength
B	1.45	1.45
D	1.61	1.31

Table IV-b
Treated Accelerated Creep Results
full log-log plot

Sheet Type	Accelerated Creep at 25% of sample's tensile strength	Accelerated Creep at 25% of B's tensile strength
B	2.26	2.26
D	2.21	2.31

Notice that comparisons of accelerated creep rates are not consistent between methods. We are therefore left in a predicament since the choice of method is at the discretion of the analyst.

DISCUSSION

We did achieve (see Table I.) a moderately higher hygroexpansion in the TMP (A) sheets than in the bleached kraft softwood (B) sheets. This gave the hoped-for opportunity to assess the role of moisture sensitivity heterogeneity in accelerated creep. We associate accelerated creep with load cycling that is driven by humidity cycling. All other things being equal, we expect the more hygroexpansive sheet to exhibit more accelerated creep. There are six (three methods at two loads) different comparisons between accelerated creep of the A samples and the B samples. In all cases, the A samples exhibited more accelerated creep. The average ratio of A to B accelerated creep is 1.23 with a standard deviation of 0.28. We can very roughly assert that A is 25% more prone to accelerated creep than is B.

In every comparison, but one, the accelerated creep numbers for the TMP-BKS blend sheets were intermediate between the pure TMP and pure BKS. The average of AB accelerated creep divided by the mean of the A and the B accelerated creep is 0.99 with a standard deviation of 0.04. Clearly, heterogeneity due to fiber type is not producing a noticeable influence on accelerated creep. This observation supports the premise that the major part of the influence (if any) of fiber level heterogeneity on cyclic moisture induced load cycling comes from fiber anisotropy rather than from fiber variety. The axial to radial difference in fiber properties dwarfs the difference between fibers. Therefore, we should not expect the overall level of load cycling to be significantly increased by the introduction of relatively small fiber-to-fiber differences.

The construction of through-sheet heterogeneity by wet pressing TMP and BKS sheets together did influence accelerated creep. In all, but one, of the nine (three methods at three loads) cases, the mixed-ply sheets had greater accelerated creep than either of the four-ply sheets made of sheets of the same furnish. The average of the sum of the mixed-ply sheets to the sum of the pure-ply sheets is 1.18 with a standard deviation of 0.15. From our perspective (5,6), the added multi-ply heterogeneity caused enough extra load cycling to perceptibly augment accelerated creep. Incidentally, the multi-ply accelerated creep values are below the single-ply numbers because of their increased sorption times. When sorption time approaches the moisture cycling time, moisture cycling (and thereby load cycling) is attenuated in the interior of the sheet (5,6).

Treatment in the Hobart mixer to attempt to microcompress the bleached kraft southern-pine fibers and hopefully to reduce anisotropy in the fiber response to moisture had little effect on accelerated creep. The average ratio of D to B accelerated creep is 0.97 with a standard deviation of 0.09. Based on our modeling (5,6), there is a maximum in accelerated creep when it is plotted as a function of fiber anisotropy. If the stiffness anisotropy is too great, most of the load will always be carried by the stiffer element, and no load cycling occurs. If the heterogeneity is too small, then elements move together,

and again there is no anisotropy-driven load cycling. It is hard to guess where on the accelerated creep vs. fiber anisotropy curve the untreated and microcompressed samples lie, and it is hard to predict which should be larger.

SUMMARY

We were able to produce handsheets of different furnishes having different hygroexpansivities and different accelerated creeps. Blends of the furnishes produced samples of intermediate hygroexpansion and accelerated creep. We therefore believe that blending of fiber types will have neither a corrupting nor a salutary influence on accelerated creep. However, we predict that multi-ply sheets of furnishes with different hygroexpansivities will experience exacerbated accelerated creep. Papermakers should consider degradation of product performance in cyclic humidity environments when they produce multi-ply products.

Acknowledgements: We thank the member companies of the Institute of Paper Science and Technology for their financial support. John Waterhouse and Derek Page made contributions to the direction of this research. Miranda Bliss made our handsheets, and Kennisha Collins did the creep testing.

LITERATURE CITED

1. Armstrong, L. D. and Kingston, R. S. T., *Nature* 185 (4716): 862 (1960).
2. Byrd, V. L., *Tappi* 55(2): 247(1972)
3. Pickett, G. J., *J. Amer. Concrete Inst.* 13(4): 257(1942)
4. Wang, J. Z., Vipul, D., Glasser, W., and Dillard, D. A., in *High Temperature and Environmental Effects on Polymeric Composites* ASTM STP 1174, C. E. Harris and T. S. Gates Eds. Amer. Soc. for Testing and Mater., Philadelphia: 186(1993).
5. Habeger, C., and Coffin, D. to be published in the April 2000 edition of the *J. Pulp Paper Sci.*
6. Coffin, D. and Habeger, C., in *Proceedings: Moisture and Creep Effects on Paper, Board, and Containers*, Grenoble, France, March 18-19, 1999
7. Selway, J. W., and Kirkpatrick J., in, *Proceedings: Cyclic Humidity Effects on Paperboard Packaging*, FPL Madison, WI: 31(1992).
8. Nanko, H., and Tada, Y., *Proceedings of the 1995 International Paper Physics Conference*: 159(1995).
9. Nanri, Y., and Uesaka, T., *Tappi* 76(6): 62(1993).
10. Uesaka, T., Moss, C., and Nanri, Y., *J. Pulp Paper Sci.* 18(1): J11(1992).
11. Salmen L., *Proceedings of the Eleventh Fundamental Research Symposium Held in Cambridge*: September 1997: 374(1997).
12. Page, D., Seth, R., Jordan, B., and Barbe, M., *Papermaking Raw Materials* (Punton, ed.), Vol. 1: (Mechanical Eng. Publns. Ltd.). 183(1985).
13. Page, D., *Pulp Paper Mag. Can.* 67(1): T2(1966).
14. Brezinski, J. P., *Tappi* 39(2): 166(1956).

**80-30% RH Accelerated Creep of Sample A
at 25% of 50% RH Breaking Load**

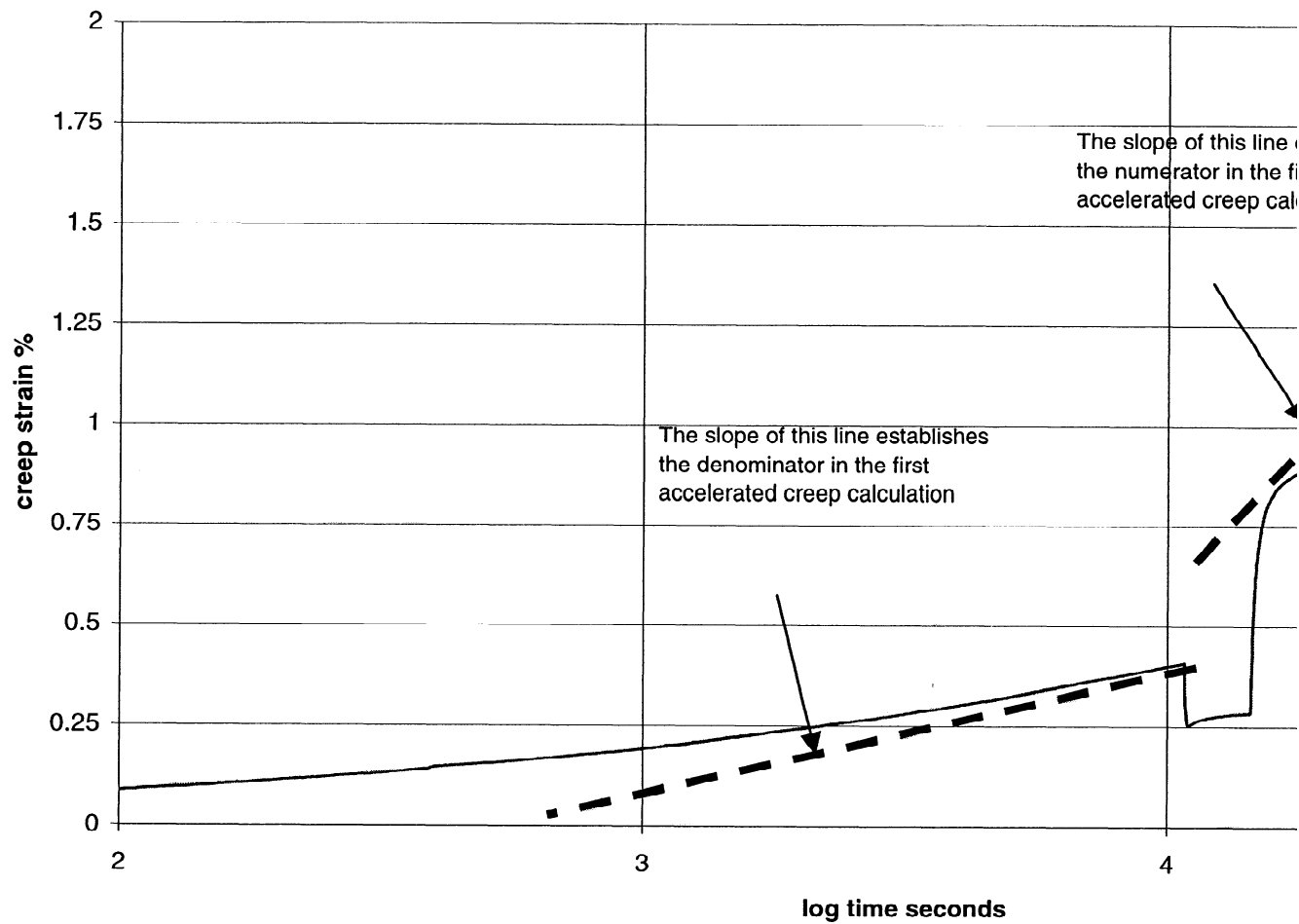


Figure 1

**80-30% RH Accelerated Creep of Sample A
at 25% of 50% RH Breaking Load**

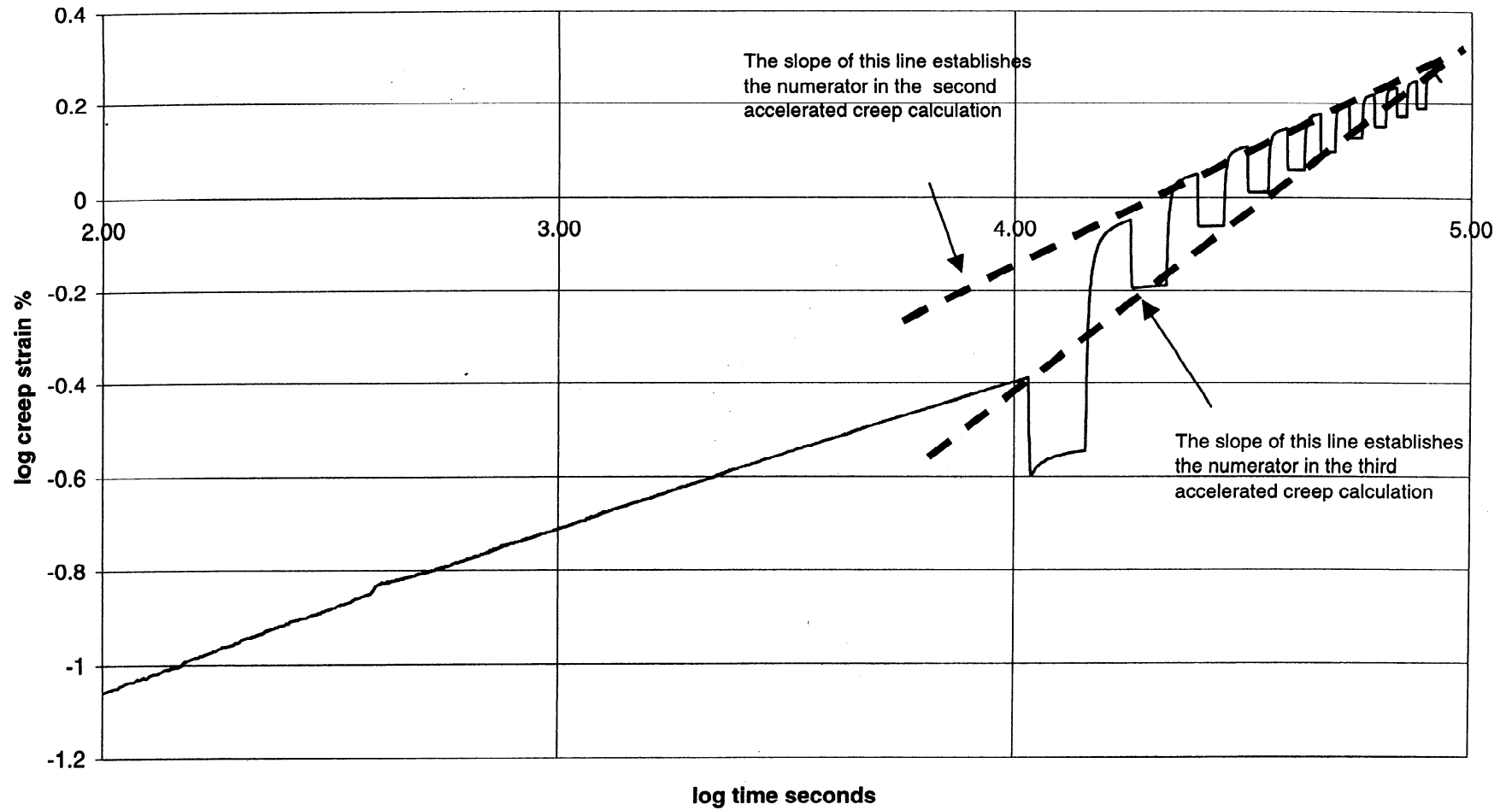


Figure 2

